



## IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

p#36

Applicant: Tadmitsu KISHIMOTO et al.

Title: PHARMACEUTICAL COMPOSITION FOR TREATMENT OF  
DISEASES CAUSED BY IL-6 PRODUCTION

RECEIVED

OCT 1-7 2002

Appl. No.: 08/817,507

TECH CENTER 1600/2900

Filing Date: 04/17/1997

Examiner: Karen A. CANELLA

Art Unit: 1642

**INFORMATION DISCLOSURE STATEMENT**  
**UNDER 37 CFR §1.56**Commissioner for Patents  
Washington, D.C. 20231

Sir:

Submitted herewith on Form PTO/SB/08 is a listing of documents known to Applicants in order to comply with Applicants' duty of disclosure pursuant to 37 CFR §1.56. A copy of each listed document is being submitted to comply with the provisions of 37 CFR §1.97 and §1.98.

The submission of any document herewith, which is not a statutory bar, is not intended as an admission that such document constitutes prior art against the claims of the present application or that such document is considered material to patentability as defined in 37 CFR §1.56(b). Applicants do not waive any rights to take any action which would be appropriate to antedate or otherwise remove as a competent reference any document which is determined to be a *prima facie* art reference against the claims of the present application.

**TIMING OF THE DISCLOSURE**

The listed documents are being submitted in compliance with 37 CFR §1.97(c), before the payment of the issue fee.

**RELEVANCE OF EACH DOCUMENT**

Any document listed on the attached PTO/SB/08 was cited as being relevant during the prosecution of the corresponding European application. A copy of the European Search Report is attached setting forth the portion of each document considered relevant by the examiner.

Applicants respectfully request that any listed document be considered by the Examiner and be made of record in the present application and that an initialed copy of Form PTO/SB/08 be returned in accordance with MPEP §609.

**STATEMENT**

The undersigned hereby states in accordance with 37 CFR §1.97(e)(1) that each item of information contained in this information disclosure statement was first cited in any communication from a foreign patent office in a counterpart foreign application not more than three (3) months prior to filing of this Statement.

The Commissioner is hereby authorized to charge any additional fees which may be required regarding this application under 37 CFR §§ 1.16-1.17, or credit any overpayment, to Deposit Account No. 19-0741. Should no proper payment be enclosed herewith, as by a check being in the wrong amount, unsigned, post-dated, otherwise improper or informal or even entirely missing, the Commissioner is authorized to charge the unpaid amount to Deposit Account No. 19-0741.

Respectfully submitted,

*Michael D. Kammori*  
By Reg. No. 32, 904

Date October 15, 2002

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Under the Paperwork Reduction Act of 1995 no persons are required to respond to a collection of information unless it contains a valid OMB control number.

**Complete if Known**

<b>Application Number</b>	08/817,507
<b>Filing Date</b>	04/17/1997
<b>First Named Inventor</b>	KISHIMOTO et al.
<b>Group Art Unit</b>	1642
<b>Examiner Name</b>	Karen A. CANELLA
<b>Attorney Docket Number</b>	053466-0201

*(use as many sheets as necessary)*

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Attorney Docket Number	053466-0201
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Examiner Initials*	Cite No. <sup>1</sup>	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.) date, page(s), volume-issue number(s), publisher, city and/or country where published.	T <sup>6</sup>
	A6	BECK et al., "Brief Report: Alleviation of Systemic Manifestations of Castleman's Disease By Monoclonal Anti-Interleukin-6 Antibody", <u>The New England Journal of Medicine</u> , March 3, 1994, Vol. 330, No. 9, pages 602-605.	
	A7	EMILIE et al., "Administration of an Anti-Interleukin-6 Monoclonal Antibody to Patients With Acquired Immunodeficiency Syndrome and Lymphoma: Effect on Lymphoma Growth and on B Clinical Symptoms", <u>Blood</u> , October 15, 1994, Vol. 84, No. 8, page 2472-2479.	

Date  
Considered

\*EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

<sup>1</sup> Unique citation designation number. <sup>2</sup>See attached Kinds of U.S. Patent Documents. <sup>3</sup>Enter Office that issued the document, by the two-letter code (WIPO Standard ST.3). <sup>4</sup>For Japanese patent documents, the indication of the year of the reign of the Emperor must precede the serial number of the patent document.

5Kind of document by the appropriate symbols as indicated on the document under WIPO Standard ST. 16 if possible. 6Applicant is to place a check mark here if English language Translation is attached.

**Burden Hour Statement:** This form is estimated to take 2.0 hours to complete. Time will vary depending upon the needs of the individual case. Any comments on the amount of time you are required to complete this form should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, Washington, D.C. 20231. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Assistant Commissioner for Patents, Washington, D.C. 20231.